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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/825,682	04/04/2001	Elena Feinstein	65503-B/JPW/MS	3555
7590	04/27/2005		EXAMINER	
John P. White Cooper & Dunham LLP 1185 Avenue Of the Americas New York, NY 10036			JOHANNSEN, DIANA B	
			ART UNIT	PAPER NUMBER
			1634	

DATE MAILED: 04/27/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

SUPPLEMENTAL EXAMINER'S AMENDMENT

1. An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

2. **Amend the specification as follows:**

Amend the specification as specified in the attached Amendment, which was requested by the examiner on March 2, 2005, and submitted to the examiner by facsimile on April 11, 2005.

3. It is noted that certain portions of the Amendment submitted April 11, 2005 are not in the format specified in MPEP 1302.04 for facsimile attachments to an examiner's amendment (specifically, 3 paragraphs provided by applicant are in a "marked-up" format, rather than a "clean" format). Accordingly, in the event that the "marked-up" versions are unusable for printing purposes, "clean" versions of the paragraphs submitted by Applicant are set forth below:

On page 55, lines 15 to 26:

--Many of the unexpected phenomena can indicate the limitation of previous understanding, and serve as a starting point for class definition. However, "outlying" hybridizations can also indicate quality problems. Overall, in each set (Figures 1-4) most of the separation between hybridizations is consistent with the expected TCC and normal urothelium separation. Even in hybridizations of lower quality, such as those of the second set, a clear separation between TCC and normal samples is observed.—

On page 55, lines 29 to 39:

--One of the TCC samples in the first set (TC6) is such an "outlyer" (Figure 1), as well as one of the normal samples in the second set, and another normal sample (TC35) in the third. (The "outlyers" do not appear to be misclassifications). For example, TC35 (a normal sample which is an "outlyer" in the third set) does not behave like a TCC sample. Rather those genes that are up-regulated in TCC samples are down-regulated in TC35).—

On page 59, lines 1 to 13:

--None of the "outlyers" was eliminated from subsequent analysis steps. Rather, they were included to facilitate the selection of a more robust marker set (Figures 1-4). Here, more complex relations are observed between global expression profiles. First, the two invasive hybridizations (TC34 and TC45), are distinct from other TCC samples (Figures 1-6). Second, the relationship between global Ta and T1 profiles is not straightforward.

Art Unit: 1634

Most of the Ta samples form a unique cluster in the 3rd set, while the T1 samples are more dispersed.—

Art Unit: 1634

4. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Diana B. Johannsen whose telephone number is 571/272-0744. The examiner can normally be reached on Monday-Friday, 7:30 am-4:00 pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, W. Gary Jones can be reached at 571/272-0745. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).



Diana B. Johannsen
Primary Examiner
Art Unit 1634
April 13, 2005

Attachment to the Supplemental Examiner's Amendment

Apr-11-05 06:59pm From-Cooper&Dunham LLP

+212 391 0526

T-398 P.001 F-408

*Requested
PPS/DS
4/13/05*

COOPER & DUNHAM LLP

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DATE : April 11, 2005 TIME: _____

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DOCKET: 2094/65503-B U.S. SERIAL NUMBER: 09/825,682
MESSAGE: AMENDMENT AFTER PAYMENT OF ISSUE FEE IN RESPONSE TO MARCH 2, 2005 TELEPHONE REQUEST BY EXAMINER JOHANNSEN

=====
THE INFORMATION CONTAINED IN THIS FACSIMILE TRANSMISSION IS INTENDED SOLELY FOR THE PERSONAL AND CONFIDENTIAL USE OF THE DESIGNATED RECIPIENT(S) NAMED ABOVE. THIS TRANSMISSION MAY BE AN ATTORNEY-CLIENT COMMUNICATION CONTAINING INFORMATION THAT IS PRIVILEGED AND CONFIDENTIAL. IF THE READER OF THIS MESSAGE IS NOT A DESIGNATED RECIPIENT OR AN AGENT RESPONSIBLE FOR DELIVERING IT TO A DESIGNATED RECIPIENT, YOU ARE HEREBY NOTIFIED THAT YOU HAVE RECEIVED THIS DOCUMENT IN ERROR, AND THAT ANY REVIEW, DISTRIBUTION, OR COPYING OF THIS MESSAGE IS STRICTLY PROHIBITED. IF YOU HAVE RECEIVED THIS COMMUNICATION IN ERROR, OR IF UPON READING THIS DOCUMENT YOU HAVE REASON TO BELIEVE THAT THE DOCUMENT WAS INADVERTENTLY SENT TO YOU, PLEASE NOTIFY US IMMEDIATELY BY COLLECT TELEPHONE CALL AND RETURN THE ORIGINAL MESSAGE TO US BY MAIL. THANK YOU.

Docket. No. 65503-B/JPW/DNS

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Elena Feinstein and Orna Mor

Serial No.: 09/825,682 Group Art Unit: 1634

Filed : April 4, 2001 Examiner: D. Johannsen

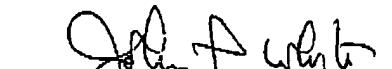
For : METHODS OF DIAGNOSING BLADDER CANCER

1185 Avenue of the Americas
New York, New York 10036
April 11, 2005BY FACSIMILE - (571)273-0744Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450FACSIMILE CERTIFICATION OF TRANSMISSION
IN CONNECTION WITH THE ABOVE-IDENTIFIED APPLICATION

Date of Facsimile: April 11, 2005. I hereby certify that this paper including Amendment After Payment of Issue Fee in Response to March 2, 2005 Telephone Request By Examiner Johannsen is being transmitted to the U.S. Patent and Trademark Office on the date indicated above by facsimile and is addressed to U.S. Patent and Trademark Office, P.O. Box 1450, Alexandria, VA 22313-1450 Attn: D. Johannsen.

Printed Name: Daniel Smith

Respectfully submitted,



John P. White
Registration No. 28,678
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1185 Avenue of the Americas
New York, New York 10036
(212) 278-0400

Docket No. 65503-B/JPW/DNS

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicants: Elena Feinstein and Orna Mor

Serial No.: 09/825,682 Examiner: D. Johannsen

Filed : April 4, 2001 Group Art Unit: 1634

For : METHODS OF DIAGNOSING OF BLADDER CANCER

1185 Avenue of the Americas
New York, New York 10036
April 11, 2005BY FACSIMILE - (571)-273-0744Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Sir:

AMENDMENT AFTER PAYMENT OF ISSUE FEE IN RESPONSE TO MARCH 2,
2005 TELEPHONE REQUEST BY EXAMINER JOHANNSEN

This Amendment is submitted in response to a March 2, 2005 telephone request from Examiner Johannsen, communicated to Daniel N. Smith of the undersigned's office during which Examiner Johannsen requested amendments to the specification of the above-identified application. Accordingly, applicants submit this Amendment in response to the Examiner's request.

Amendments to the Specification begin on page 2 of this paper.

Remarks/Arguments begin on page 32 of this paper.

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Serial No.: 09/825,682
Filed: April 4, 2001
Page 2

Amendments to the Specification

Please add the following section on page 8, line 5 of the specification immediately before the beginning of "DETAILED DESCRIPTION OF THE INVENTION":

-- BRIEF DESCRIPTION OF THE FIGURES

Figure 1: This figure shows relationships between hybridizations' hierarchical clustering of Set #1 hybridizations. Either the Pearson correlation coefficient or a standard Euclidean distance was used as the distance measure between differential hybridization vectors. Hybridizations were clustered according to these distances by average linkage hierarchical clustering. Missing values were deleted on a case-wise basis. Clusters of Missing values hybridizations can be identified and evaluated in light of existing knowledge.

Figure 2: This figure shows relationships between hybridizations' hierarchical clustering of Set #2 hybridizations. Either the Pearson correlation coefficient or a standard Euclidean distance was used as the distance measure between differential hybridization vectors. Hybridizations were clustered according to these distances by average linkage hierarchical clustering. Missing values were deleted on a case-wise basis. Clusters of Missing values hybridizations can be identified and evaluated in light of existing knowledge.

Figure 3: This figure shows relationships between hybridizations' hierarchical clustering of Set #3 hybridizations. The figure is a tree diagram for 21 variables with unweighted pair-group average. The Pearson correlation coefficient was used as the distance measure between differential hybridization vectors. Hybridizations

Applicants: Elena Feinstein and Orna Mor
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Page 3

were clustered according to these distances by average linkage hierarchical clustering. Missing values were deleted on a case-wise basis. Clusters of Missing values hybridizations can be identified and evaluated in light of existing knowledge.

Figure 4: This figure shows relationships between hybridizations' hierarchical clustering of Set #3 hybridizations. The figure is a tree diagram for 21 variables with unweighted pair-group average. The standard Euclidean distance was used as the distance measure between differential hybridization vectors. Hybridizations were clustered according to these distances by average linkage hierarchical clustering. Missing values were deleted on a case-wise basis. Clusters of Missing values hybridizations can be identified and evaluated in light of existing knowledge.

Figure 5: This figure shows relationships between hybridizations' hierarchical clustering for all TCC sample hybridizations. The figure includes raw data with unweighted pair-group average. The Pearson correlation coefficient was used as the distance measure between differential hybridization vectors. Hybridizations were clustered according to these distances by average linkage hierarchical clustering. Missing values were deleted on a case-wise basis. Clusters of Missing values hybridizations can be identified and evaluated in light of existing knowledge.

Figure 6: This figure shows relationships between hybridizations' hierarchical clustering for all TCC sample hybridizations. The figure includes raw data with unweighted pair-group average. The standard Euclidean distance was used as the distance measure between

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Filed: April 4, 2001
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differential hybridization vectors. Hybridizations were clustered according to these distances by average linkage hierarchical clustering. Missing values were deleted on a case-wise basis. Clusters of Missing values hybridizations can be identified and evaluated in light of existing knowledge.--

Please amend the paragraph on page 55, lines 15 to 26 of the specification as follows:

--Many of the unexpected phenomena can indicate the limitation of previous understanding, and serve as a starting point for class definition. However, "outlying" hybridizations can also indicate quality problems. Overall, in each set ~~(Table B3)~~ (Figures 1-4) most of the separation between hybridizations is consistent with the expected TCC and normal urothelium separation. Even in hybridizations of lower quality, such as those of the second set, a clear separation between TCC and normal samples is observed.--

Please amend the paragraph beginning on page 55, lines 29 to 39 of the specification as follows:

-- One of the TCC samples in the first set (TC6) is such an "outlyer" ~~(Table B3)~~ (Figure 1), as well as one of the normal samples in the second set, and another normal sample (TC35) in the third. (The "outlyers" do not appear to be misclassifications). For example, TC35 (a normal sample which is an "outlyer" in the third set) does not behave like a TCC sample. Rather those genes that are up-regulated in TCC samples are down-regulated in TC35).--

Please delete pages 56 to 58 of the specification.

Applicants: Elena Feinstein and Orna Mor
Serial No.: 09/825,682
Filed: April 4, 2001
Page 5

Please amend the paragraph beginning on page 59, lines 1 to 13 of the specification as follows:

-- None of the "outlyers" was eliminated from subsequent analysis steps. Rather, they were included to facilitate the selection of a more robust marker set (~~Table B3~~) (Figures 1-4). Here, more complex relations are observed between global expression profiles. First, the two invasive hybridizations (TC34 and TC45), are distinct from other TCC samples (~~Tables B3 and B4~~) (Figures 1-6). Second, the relationship between global Ta and T1 profiles is not straightforward. Most of the Ta samples form a unique cluster in the 3rd set, while the T1 samples are more dispersed.--

Please renumber pages 59 to 82 of the specification as pages 56 to 79.

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 Page 6

Please replace the existing Table E on pages 83 and 84 of the specification with the following new Table E expanded into 8 pages to improve legibility:

TABLE E (Page 1)

Index	GeneDescription
1	Human 54 kDa protein mRNA, complete cds ; nt non_genomic(Identity) REPEAT:contig ICC_29E6_RF.fa
2	none;1 ICC_74D3_1_M13F.fa none;25 ICC_74D3_1_M13R.fa
(A)121586) d477C04_1 (novel protein similar to orofillin and dystrophin) (Homo sapiens); nt(non_genomic(Identity))10 ICC_70E2_1_M13F.fa Homo sapiens serine protease inhibitor, Kunlitz type, 2 (SPIN12), mRNA;	
3	nt(non_genomic(Identity))24 ICC_70E2_1_M13R.fa
4	Homo sapiens full length insert cDNA clone ZC48G12 ; nt non_genomic(Identity):contig ICC_75B7_RF.fa
5	nt non_genomic(Identity):contig ICC_89H5_RF.fa
6	Homo sapiens cDNA, FLIP2720 fs, clone HS14320 nt non_genomic(Identity);16 ICC_10D4_1_M13F fa ANKHZN protein [Homo sapiens]300.1 (AB037360) ANKHZN (Homo sapiens); nt(non_genomic(Identity))30 ICC_10D4_1_M13R.fa low molecular mass ubiquinone-binding protein (Homo sapiens) UCRQ_HUMAN_UBIQUINOL-CYTOCHROME C REDUC; nt(non_genomic(Identity))contig ICC_67C11_RF.fa
7	none;14 ICC_79B1_1_M13F fa Homo sapiens quiescin Q6 (QSCN6), mRNA;
8	nt non_genomic(Identity):contig ICC_79B1_1_M13R.fa
9	Homo sapiens mRNA, cDNA, DKFZp564C2282 (from clone DKFZp564C2282);
10	Homo sapiens full length insert cDNA clone ZC48G12 ; nt non_genomic(Identity):contig ICC_5&G1_RF.fa
11	NADH dehydrogenase (ubiquinone) 1, subcomplex unknown, 2 (14.5k, B14.5b) (Homo sapiens) N4BM_HUMAN ; nt(non_genomic(Identity))contig ICC_60B2_RF.fa
12	
13	
14	

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TABLE E (Page 2)

Acc1	GenID	TC7A_BDE_DIFF	TC8A_BDE_DIFF	TC9A_BDE_DIFF	TC10A_BDE_DIFF
gl 407307 gb U02493.1 HSU02493	ICC-29E_6	-1	-1.2	1.1	1
	ICC-74D_3	-1.1	-1	1.2	-1.1
gl 7671662 emb CABB9410.1					
gl 10863908 ref NM_021102.1	ICC-70E_2	-1.2	1.1	1.2	-1.1
gl 3483555 gb AF086210.1 HUMZC48G12	ICC-75B_7	-1.1	1	1.2	-1.2
gl 7656943 ref NM_014400.1	ICC-89H_5	-1.1	-1	1.1	1
gl 10439219 db AK026373.1 AK026373	ICC-101D_4	-1.1	-1.1	1.1	-1.1
gl 17705278 ref NP_057460.1					
gl 7657486 ref NP_055217.1	ICC-67C_11	1	-1.1	-1	1
gl 4506360 ref NM_002826.1	ICC-798_1	-1.1	1	12	-1.1
gl 7328021 emb AU101965.1 HSMA02543	ICC-74H_12	-1.1	-1.1	13	-1.1
gl 3483555 gb AF086210.1 HUMZC48G12	ICC-56G_1	-1	-1	13	-1.1
	ICC-47D_9	1.1	1	-1	-1.1
gl 4758784 ref NP_004640.1	ICC-60B_2		-1.1	-1.1	-1.1
	ICC-23D_4	1.1	-1.4	-1.2	1.2
	ICC-20F_5	-1.2	1.1	12	-1.2

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TABLE 3 (Page 3)

TC11A_BDE_DIFF	TC22A_BDE_DIFF	TC21A_BDE_DIFF	TC23A_BDE_DIFF	TC24A_BDE_DIFF	TC47A_BDE_DIFF	TC39A_BDE_DIFF
1.2	-1.1	1.3	-1.2	1+	-1	1.1
-1	1.1	1	1	-1.2	1	1
-1.1	1.2	1.1	1	-1.4	1.2	-1
-1	1.1	-1	1	-1.2	1.1	1
-1	1.2	-1	1.1	-1.2	1.1	1
1.2	1.1	1.1	1.1	-1.3	1.1	-1.1
1.1	1.1	1.1	-1	-1.2	-1	-1
-1	1.1	-1	1.1	-1.2	1.1	-1
1	1.2	1.2	-1	-1.4	1.1	-1.3
-1.1	1.2	1	1	-1.2	-1	1.1
-1	-1.3	1.4	-1.1	1+	-1.1	-1
13	-1	1.2	-1.1	1+	1.1	-1.2
13	-1.2	1.1	1.1	1+	-1.1	1.1
1.1	1	1.1	-1.2	1		-1.2

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TABLE 2 (Page 4)

TC38A_BDE_DIFF	TC46A_BDE_DIFF	TC48A_BDE_DIFF	TC37A_BDE_DIFF	TC35A_BDE_DIFF	TC2A_BDE_DIFF	TC3A_BDE_DIFF
-1,1	1,1	1,4	-1	-1,5	1,1	-1,2
1,3	1,3	1,1	-1	-1,8	-1,5	-1,1
1,3	1,3	1,2	-1,1	-2,1	-1,3	-1,5
1,3	1,3	1,1	-1	-1,9	-1,2	1
1,3	1,3	1,1	-1,1	-1,9	-1,3	-1,1
1,2	1,3	1,1	-1,1	-1,7	-1,2	-1,1
1,2	1,1	1	-1,2	-1,2	-1,4	1,1
1,2	1,3	1,1	-1	-1,8	-1,2	-1
1,3	1,3	1,2	-1,1	-1,6	-1,2	-1,3
1,3	1,3	1,1	-1,1	-1,8	-1,4	-1
-1,1	1,2	1,3	1,2	-1,7	-1,2	-1,2
-1,1	1,3	1,1	-1,1	-1,2	1	1,2
-1	1,1	1,3	1,1	-1,7	-1,1	-1,2
1,1	1,1	1,1	-1	-1,3	-1	1

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TABLE E (Page 5)

TC4A_BDE_DIFF	TC5A_BDE_DIFF	TC8A_BDE_DIFF	TC16A_BDE_DIFF	TC17A_BDE_DIFF	TC18A_BDE_DIFF	TC25A_BDE_DIFF
1.4	1.2	-1.1	-1	-1.5	-1	1.4
1	1.5	1.1	1	1.3	1.5	1.5
-1.3	1.3	-1	1.2	1.5	1.3	1.4
-1.4	1.3	1.3	1.1	1.4	1.3	1.5
-1.3	1.2	1	1.1	1.5	1.1	1.5
-1.5	-1	-1	1	1.4	1.7	1.5
1.1	-1	-1	1.1	1.5	1.5	1.5
1.1	-1.2	1	1	1.4	1.4	1.5
-1.1	1.1	-1.2	1.1	1.4	1.4	1.5
-1.4	1.3	1.2	1.1	1.4	1.5	1.5
-1.6	1.2	-1	1.1	-1.3	1	1.3
-1.2	1	1	1.4	1.2	1.3	1.5
-2	1.4	1.1	-1.4	-1.5	-1.3	1.2
-1.2	-1.1	-1.2	1.2	1.4	1.3	1.3

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TABLE E (Page 6)

TC19A_BDE_DIFF	TC20A_BDE_DIFF	TC32A_BDE_DIFF	TC33A_BDE_DIFF	TC41A_BDE_DIFF	TC42A_BDE_DIFF	TC44A_BDE_DIFF
1.1	1.4	-1.4	-1.1	-1.1	-1.3	-1.3
1.1	1.3	-1.2	1.1	1.1	1.1	1.2
1.2	1.3	-1.3	-1.2	1.2	-1	1.2
1.4	1.1	-1.1	-1	1.1	1.1	1.1
1.4	1.1	-1.1	-1	1.1	1	-1.1
12	13	-1.3	-1.2	1.1	-1.1	-1
1.4	1.4	-1.3	-1.3	1.1	-1	-1
1.4	1.1	-1.1	-1	1.1	-1	1.1
1.2	1.4	-1.5	-1.4	1.1	-1.1	1.2
1.4	1.3	-1.1	1	1.2	1	1.1
13	1.2	-1.5	-1.2	-1.2	-1.3	-1.3
1.2	1.2	-1.3	-1.2	-1.1	-1.4	-1.2
-1	-1.1	-1	-1.1	1	-1.3	-1.1
1.4	1.1	-1.2	-1.5	-1	-1.2	-1.3

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TABLE E (Page 7)

TC31A_BDE_DIFF	TC39A_BDE_DIFF	TC43A_BDE_DIFF	TC40A_BDE_DIFF	TC30A_BDE_DIFF	TC28A_BDE_DIFF	TC29A_BDE_DIFF
-1.1	-1.2	-1.3	-1.3	-1.3	1.2	1.2
-1.3	-1.1	-1.4	-1.2	1.1	-1.1	1.2
-1.3	-1.1	-1.4	-1.2	1.1	-1.1	1.1
-1.3	-1.1	-1.5	-1.3	1.1	-1	1.1
-1.2	-1.2	-1.5	-1.2	1.1	-1.1	1.1
-1.3	-1.2	-1.4	-1.4	1.1	-1.3	-1.1
-1.4	-1.2	-1.4	-1.5	-1	-1.1	-1
-1.1	-1.1	-1.4	-1.3	1.1	-1.1	1.2
-1.3	-1.2	-1.5	-1.4	1.2	-1.1	1
-1.3	-1.1	-1.5	-1.3	1	-1.1	1.1
-1	-1.4	-1.4	-1.5	-1.5	1.2	1.2
-1.2	-1.9	-1.4	-1.3	-1.3	-1.1	-1.2
12	-1.3	1	-1.1	-1.2	1.4	1.4
-1.3	1	-1.2	-1	-1.3	1.1	-1.2

Applicants: Elena Feinstein and Orna Mor
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Page 13

TC34A_BDE_DIFF	TC45A_BDE_DIFF
1.1	1.1
1.4	1.1
13	1.2
12	1.1
1.2	-1
1.1	1.1
-1.1	-1
1.4	1.1
1	1.3
1.3	1
1.1	-1.1
12	1.1
1.3	-1.1
-1.2	1.2

TABLE E (Page 8)

- -

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Please replace the existing Table 1 on pages 85 and 86 of the specification with the following new Table 1 expanded into 8 pages to improve legibility:

TABLE 1 (Page 1)

GeneDescription and clone ID
1 Human sepiolite keratin 19 (KRT19) gene, complete cd ; nt_non_genomic[identity]47_TCC_91B11_M13F.TXTfa
2 Human 40-kDa keratin intermediate filament precursor; nt_non_genomic[identity]20_TCC_60H4_M13F.TXTfa
3 Human mRNA fragment for mesothelial type II kerat ; nt_non_genomic[identity]04_TCC_94G3_M13F.TXTfa
4 Homo sapiens cystatin B (CSTB) gene, promoter reg ; nt_non_genomic[identity]30_TCC_76B3_M13F_F04_042.ab1.TXT
5 Homo sapiens S100 calcium-binding protein P (S100P ; nt_non_genomic[identity]29_TCC_48G1_M13F.TXTfa
6 Homo sapiens syntetan 1 (SDCT1) mRNA ; nt_non_genomic[identity]26_TCC_75E3_M13F_804_032.ab1.TXT
7 Homo sapiens S100 calcium-binding protein A13 (S10 : nt_non_genomic[identity]28_TCC_44C1_M13F.TXTfa
8 Homo sapiens mRNA for hepatocyte growth factor ac ; nt_non_genomic[identity]35_TCC_21D6_M13F_C05_037.ab1.fa
9 Homo sapiens midkine [neutile growth-promoting fac ; nt_non_genomic[identity]32_TCC_89G3_M13F_811_092.ab1.TXT
10 Homo sapiens solute carrier family 2 (facilitated ; nt_non_genomic[identity]37_TCC_57B3_M13F.TXT
11 Homo sapiens S100 calcium-binding protein A11 (cal ; nt_non_genomic[identity]31_TCC_65B9_M13F.TXTfa
12 Homo sapiens fatty acid binding protein 5 (fasfam ; nt_non_genomic[identity]11_TCC_25F2_M13F.TXT
13 Homo sapiens Oprm1 interacting protein OIP3 mRNA, p ; nt_non_genomic[identity]38_TCC_56E11_M13F.TXT
14 Homo sapiens glutaminyl-tRNA synthetase (GARS), m ; nt_non_genomic[identity]46_TCC_78B11_M13F_506_058.ab1.TXT
15 Homo sapiens anterior gradient 2 (Xenopus laevis) ; nt_non_genomic[identity]25_TCC_50G5_M13F.TXT
16 Homo sapiens myristoylated alanine-rich protein k ; nt_non_genomic[identity]10_TCC_53H11_T3
17 Homo sapiens autophagy-associated phosphoprotein pf ; nt_non_genomic[identity]46_TCC_27H5_M13F_F08_058.ab1.fa
18 Homo sapiens type II membrane serine protease (LLO ; nt_non_genomic[identity]53_TCC_78G2_M13_F007_054.ab1.TXT
19 Homo sapiens putative secreted protein XAG mRNA, c ; nt_non_genomic[identity]26_TCC_50G8_M13F.TXT
20 H.sapiens (ssca) mRNA, 340bp ; nt_non_genomic[identity]40_TCC_1SF1_L_M13F.fa

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TABLE 1 (Page 2)

	G2/3	G2/3
	T1	T1
Accession	GeneID	TCC_A_BDE_DIFF
g 6729880 gb AF202321.1 AF202321	TCC-91B_11	3.7
g 184688 gb J03607.1 HUMIFP	TCC-60H_4	3.7
g 34067 amb X03212.1 HSKE77R	TCC-91G_3	3.3
g 7263011 gb AF208234.1 AF208234	TCC-76B_3	5.7
g 5174662 ref NM_005980.1	TCC-48G_1	2.7
g 4506858 ref NM_002897.1	TCC-75E_3	1.9
g 5174658 ref NM_005979.1	TCC-44C_1	3.1
g 2924619 gb AB006534.1 AB006534	TCC-21D_6	1.1
g 4505134 ref NM_002391.1	TCC-89G_3	2.2
g 5730050 ref NM_008516.1	TCC-57B_3	4.8
g 5032056 ref NM_008820.1	TCC-68B_9	3.5
g 4557580 ref NM_001444.1	TCC-25F_2	2.1
g 2815605 gb AF025439.1 AF025439	TCC-56E_11	3.4
g 4826865 gb AF025439.1 AF025439	TCC-78B_11	2.2
g 5453540 ref NM_008408.1	TCC-50G_5	2.4
g 4505082 ref NM_002356.1	TCC-53H_11	1.8
g 5031650 ref NM_005563.1	TCC-27H_5	1.8
g 705976 ref NM_016425.1	TCC-79G_2	5.6
g 6652811 gb AF088867.1 AF088867	TCC-50G_6	2.3
g 533886 emb Z36852.1 HSXSCAD	TCC-13F_11	-1.2
		-1.1
		-2

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TABLE I (Page 3)

G2/3	G1/2							
T1	T1	[normal]	normal	normal	normal	normal	normal	normal
TC5A_BDE_DIFF	TC6A_BDE_DIFF	TC7A_BDE_DIFF	TC8A_BDE_DIFF	TC9A_BDE_DIFF	TC10A_BDE_DIFF	TC11A		
1.5	3.1	1	-2.5	-2	1.2	-1.2		
1.5	2.9	1	-2.6	-1.9	1.3	-1.1		
3.1	1.6	1	-2.1	-1.8	1.3	-1.4		
1.6	1.3	1	-2	-1.8	-1	-1.2		
1.7	1.7	1	-2.6	-2.5	2.1	1.6		
1	1.6	1	-1.7	-1.3	-1.1	-1.5		
1.8	1.8	1	-2.8	-2.4	1.9	1.4		
2	1.8	1	-1.6	-1.2	1.1	-1.5		
-1.5	1.9	1	-1.2	1.2	-1.6	-1.2		
1.2	3.2	1	-2.1	-1.5	1.1	-1.6		
1.5	1.6	1	-1.4	-1.7	1.1	-1.1		
2.6	5	1	-1.5	-1.5	2	1		
1.1	1.6	1	-1.1	-1.1	-1.3	1.1		
1.8	1.6	1	-1.1	-1.3	1.3	1.6	1.1	
-1.2	1.2	1	-2	-1.9	1.5	1.2		
1.6	1.7	1	-1.3	1.3	1.6	1.1		
2.2	1.6	1	-1.2	1.2	-1.1	1.1		
-1.6	-1.2	1	-2.2	-2.1	1.1	-1.2		
-1.1	1.3	1	-1.8	-1.9	1.4	1.2		
1.8	1.8	1	-1.6	-1.3	-1	-1.5		

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TABLE 1 (Page 4)

G2	G2	High	Low	G3
T1	Ta	Ta	T1	T1
TC16A_BDE_DIFF	TC17A_BDE_DIFF	TC18A_BDE_DIFF	TC19A_BDE_DIFF	TC20A_BDE_DIFF
2.6	2.4	2.5	2.6	1.7
2.4	2.6	3.6	2.9	1.6
7.1	4.6	6.3	4.1	4.3
3.6	11	8.9	2.3	3.4
3	6.1	8.2	2.8	1.3
5	4.6	4	2.4	4.7
2.8	4.9	6.8	2.5	1.1
2.7	2.8	2.6	1.7	2
1.9	2.5	1.9	1.4	2.9
2.7	1.8	2.9	-1	1.4
2.9	3.8	2.7	2.3	1.5
1.2	1.7	4.6	2.2	1.6
2.8	3.6	4.5	2.3	1.7
2.7	5	2.1	1.9	1.8
2.8	2.8	2.9	4.9	1.6
1.9	1.6	2.5	1.5	2.2
-1.1	-1.4	1.4	-1.4	1.2
3.8	1.4	3.6	3.9	-1.2
3.1	3.7	4.5	4.4	2.1
1.8	2.1	2.2	1.2	1.8
				-1.3

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TABLE 1 (Page 5)

normal	normal	normal	T1	T1+TIS	T1	High	G3
TC22A_BDE_DIFF	TC23A_BDE_DIFF	TC24A_BDE_DIFF	TC25A	TC28A_BDE_DIFF	TC29A_BDE_DIFF	T1	G3
1	1.3	-2.1	2.6	2.4	2.2	2	
1	1.2	-1.9	2.1	2.2	2	1.8	
1	1.8	-1.7	4	1.5	3.5	3.6	
1	1.7	-1.8	4.4	-1.3	2	2.1	
1	1.9	-1.1	4.4	-1.1	3.5	3.2	
1	1.4	-1.1	2.7	1.6	1.2	1.3	
1	1.9	-1.1	3.5	1.2	2.4	2	
1	1.5	1.7	2.4	3.9	2.2	1.7	
1	1.4	1	3.7	7	2	-1.1	
1	1.2	-1.2	2.4	-1.1	2	2.7	
1	1.2	-2.4	3.8	-1.1	2.4	1.1	
1	1.1	1	1.8	-1.8	1	5.7	
1	1.4	-1.2	3.1	-1.3	1.2	1.1	
1	1.2	-2	2.4	1.1	1.7	1.3	
1	1.2	1.4	3.1	-1.3	1.8	1.2	
1	1.1	-2.3	3.9	-1.3	2.5	2	
1	1.1	1.6	-1.3	5.7	6.7	1.9	
1	2.1	1.4	3.8	-2.1	-1.5	1.5	
1	1.5	1.2	3.7	-1.5	1.5	-1	
1	-1.1	-1.3	1.8	4.6	2.8	1.9	

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G1/2	G2	G2			INV	normal	normal
Ta	Ta/T1	Ta					
TC31A_BDE_DIFF	TC32A_BDE_DIFF	TC33A_BDE_DIFF	TC34A_BDE_DIFF	TC35A_BDE_DIFF	TC36A_BDE_DIFF		
2.1	2.3	2.4	-1	-3.9	-1.4		
1.8	2	1.9	-1.2	-4.5	-1.6		
4.3	3.2	2.1	1.1	-3.3	-1.2		
1.6	4.8	2	3.3	-1.4	-1		
3.7	5	2.6	1.8	-4	1		
4.3	3.7	2.2	1.1	1.1	-1		
2.4	3.1	1.7	1.6	1.2	1		
2.5	2.8	2.7	2.1	2	-1.3		
2.5	1.8	2.5	1.3	-1.3	1.1		
13	1.8	3.8	3	1.4	1.1		
2.3	2.4	1.8	2.5	-1.5	-1.2		
3.5	1	3.9	-1.1	-4.1	-1.5		
3.1	1.6	1.8	-1.1	2	-1.2		
-1	2	1.9	-1.1	-2.3	-1.4		
2.3	6.7	-1.3	1.3	-1.3	1.2		
3.7	2.8	2.5	2.1	-1.2	1.1		
1.8	1	1.7	3.5	1.5	1.3		
5.3	-1.9	3.2	-1.8	-1.8	-1.2		
2.1	6	-1.5	1.1	-1.3	-1.1		
3	3.1	3.1	2.5	2.4	-1.1		

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TABLE 1 (Page 7)

	Low	G2/3	G2	G2
normal	normal	T _a	T _a	T _a
TC37A_BDE_DIFF	TC38A_BDE_DIFF	TC39A_BDE_DIFF	TC40A_BDE_DIFF	TC41A_BDE_DIFF
-1.8	-1.4	3.2	3.1	2.2
-2.3	-1.8	2.9	2.5	1.8
-1.4	-1.3	3.5	-1.0	2.4
-1.8	-1	4.1	2.3	1.7
-1.7	-1.3	3.4	4.5	1.3
-1	1	2.3	3.4	2
-1.6	-1	2.4	2.7	1.1
-1.8	1	1.4	1.6	2.6
-1.2	1	1.8	4	1.7
-1.4	1.1	3.8	3.4	3.2
-1.8	-1.1	2.7	1.2	2.1
-1.9	-1.1	1.1	2.8	2.5
-1.4	-1	2.3	2	1.8
-1.8	-1.3	1.8	2.6	1.1
-1.3	1.1	1.9	1.5	1.9
-1.2	1.1	1.3	1.6	2
-1.2	1.1	1.9	2.5	2.4
-1.1	-1.1	2.2	1.5	2.3
1	1	1.8	1.2	1.9
-1.7	1.2	1.7	1.9	2.9
				2

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TABLE 1 (Page 8)

G2	G2							
T1	Ta	INV	normal	normal	normal	normal	normal	secreted
TC43A_BDE_DIFF	TC44A_BDE_DIFF	TC45A_BDE_DIFF	TC46A_BDE_DIFF	TC47A_BDE_DIFF	TC48A_BDE_DIFF	TC49A_BDE_DIFF	TC50A_BDE_DIFF	
1.7	3.6	1.1	-1	1	-1.1	3.9		
1.6	3.6	1.1	-1.1	1	-1.1	3.8		
1.8	5.6	1.3	-1.1	1	-1.2	3.8		
2.2	3.3	1.2	1.1	1	-1.1	-1.8		
2.3	3.4	1.6	-1.2	1	-1	-1.2		
2.2	3.1	1.1	-1	1	-1.1	1		
1.5	2.9	-1.4	1.1	1	-1	-1.4		
2	3	1.2	-1.1	1	-1.3	-2.2		
1.4	2.3	1.5	1.3	1	1.1	-2.8		
2.6	3.8	2.7	1.1	1	1			
1.2	3.1	1.3	1.1	1	-1.3	-1.8		
2.2	3.9	2.1	1.1	1	-1.2	-1.2		
1.3	2.8	1.4	1	1	-1.2	4.9		
1.3	1.4	-1.3	1.1	1	-1.1	-1.5		
1.8	2.3	1.1	-1.4	1	-1.2			
2	1.3	1.4	1.6	1	-1.1	3.8		
1.4	1	2.5	1	1	1	1.9		
2.8	3.1	1.1	-1.2	1	-1.2	-1.7		
1.9	2.9	1.1	-1.3	1	-1.1	-1.4		
2.1	3.2	1.3	1	1	-1.2	-2.2		

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Please replace the existing Table 2 on pages 87 and 88 of the specification with the following new Table 2 expanded into 10 pages to improve legibility:

GeneDescription and clone ID
1 Homo sapiens full length insert cDNA clone ZC4BG12 ; nt_non_genomic[identity]:03_TCC_57E11_T7.TXT.fa
2 Xq0fh07.x1 Soares_NHCeC_cervical_tumor Homo sapien ; est[identity]:24_TCC_9BC7_M13F.TXT
3 hh68c04.x1 NCI_CGAP_GU1 Homo sapiens cDNA clone IM ; est[identity]:26_TCC_13H10_M13F_B04_032.ab1.fa
4 none:21_TCC_43E2_M13F.TXT
5 RCG-BN0053-170200-011-e12BN0053 Homo sapiens cDNA ; est[identity]:28_TCC_16D12_M13F_D04_041.ab1.fa
6 none:08_TCC_70E7_M13F_H01_015.ab1.TXT
7 Homo sapiens clone RP4-584D14, complete sequence ; nt_non_genomic[identity]:14_TCC_9B6_M13F_F02_026.ab1.fa
8 Homo sapiens CG1-81 protein (LOC51108), mRNA ; nt_non_genomic[identity]:11_TCC_10E11_M13F.TXT.fa
9 Homo sapiens mRNA for KIAA0860 protein, complete ; nt_non_genomic[identity]:72_TCC_37E11_M13F_H09_079.ab1.fa
10 Homo sapiens GaINAc-T1 gene, 3'UTR ; nt_non_genomic[identity]:54_TCC_30E5_M13F_F07_062.ab1.fa
11 none:15_TCC_71H8_M13F_G02_019.ab1.TXT
12 Homo sapiens ETAA16 protein (ETAA16), mRNA ; nt_non_genomic[identity]:09_TCC_10C11_M13F.TXT.fa
13 Homo sapiens cDNA FLJ10861, clone NT2RP400157 ; nt_non_genomic[identity]:24_TCC_12F3_M13F_H03_031.ab1.fa
14 hh75c10.x1 NCL_CGAP_GU1 Homo sapiens cDNA clone IM ; est[identity]:59_TCC_34D5_M13F_C08_065.ab1.fa
15 none:15_TCC_57C3_M13F.TXT.fa
16 YBfa11.1 Soares breast2NbHBst Homo sapiens cDNA ; est[identity]:29_TCC_17A5_M13F_E04_034.ab1.fa
17 hypothetical protein C50F7.2 - Caenorhabditis eleg : CONTCG_nristong:31_TCC_10E8_M13F.fa
18 none:13_TCC_71E4_M13F_E02_018.ab1.TXT
19 Homo sapiens mRNA; cDNA DKFZp434L0310 [from clone ; nt_non_genomic[identity]:57_TCC_80C9_M13F_A08_056.ab1.TXT
20 none:44_TCC_70EB_M13F.TXT.fa

TABLE 2 (Page 1)

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TABLE 2 (Page 2)

		T ₁	G2/3
Accession	GenoID	TC2A_BDE_DIFF	T1
gi 3483555 gb AF086210.1 HUMZC48G12	TCC-57E_11	3.3	2.8
gi 7154678 gb AW516596.1 AW516596	TCC-96C_7	2.8	4.3
gi 7318401 gb AW613215.1 AW613215	TCC-13H_10	2.2	1.6
gi 8257938 gb AW997704.1 AW997704	TCC-43E_2	2.7	4.7
gi 8468933 gb AC005586.2 AC005586	TCC-70E_7	3.8	2.1
gi 7705788 ref NM_016025.1	TCC-9B_6	11	-1
gi 3327133 gb AB014560.1 AB014560	TCC-101E_11	5.1	2.1
gi 2292903 emb Y10343.1 HSY10343	TCC-37E_11	8	-1.5
gi 9506580 ref NM_019002.1	TCC-30E_5	3.3	2.3
gi 7023162 gb AK001723.1 AK001723	TCC-71H_8	1.7	1.4
gi 7318796 gb AW613610.1 AW613610	TCC-101C_11	6.1	2.2
gi 843837 gb R70320.1 R70320	TCC-12F_3	1.4	2
gi 7497781 pir T29299	TCC-34D_5	1.6	2.5
gi 680833 emb AL137591.1 HSM802346	TCC-57C_3	3.5	2.8
	TCC-17A_5	2.7	2.6
	TCC-10E_8	1.1	1.4
	TCC-71E_4	1.7	1.6
	TCC-80C_9	3.7	2
	TCC-70E_8	4.1	2.2

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TABLE 2 (Page 3)

G2/3	G2/3	G1/2	
T1	T1	normal	normal
TC4A_BDE_DIFF	TC5A_BDE_DIFF	TC6A_BDE_DIFF	TC7A_BDE_DIFF
3.2	1.3	2.3	1
5.8	5.9	-1.4	1
2.1	1	1.2	1
7.3	1.5	1.7	1
2.4	8.8	-1.4	1
3.8	1.6	1.5	1
3.9	4.1	-1.6	1
3.1	1.6	1.3	1
3.1	3.1	-1.3	1
-1.2	1.3	2	1
-1	2.5	2	1
3.2	1.9	1.2	1
1.1	-1.3	1.6	1
1.9	1.1	1.4	1
2.9	1.2	2.3	1
1.9	1.7	9.3	1
1.6	1.5	-1.1	1
1.3	2.3	2.1	1
3.2	1.7	1.3	1
3.7	1.7	1.5	1

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TABLE 2 (Page 4)

		G2	G2	High
normal	normal	normal	Ta	Ta
TC9A_BDE_DIFF	TC10A_BDE_DIFF	TC11A	TC16A_BDE_DIFF	TC17A_BDE_DIFF
-1.5	1.2	-1.2	2.2	2.6
-1	2.7	1.2	4	7.5
1.1	1	-1	1.3	1.2
-2.3	2.1	1.7	1.8	5.1
1.3	1.3	-1.2	3.6	4.4
-2.1	1.1	-1.1	3.5	4.8
1	-1.2	-1.4	3	3.7
-1.4	1.1	1.2	2.8	7.2
-1.2	-1.4	-1.6	1.6	2
1.1	1.8	2.1	-1.1	1.7
-1.2	1.1	-1.3	2.1	2.6
-1.4	-1	1	2.6	6.8
1.1	2	1.2	1.2	1.7
1.1	1.9	1.4	-1.7	-1.2
-1.6	1.2	1.2	1.6	2.6
1.5	1.8	1.6	1.5	-1
1.3	1	-1.1	-1.1	-1.7
-1.3	1	-1.1	1.8	2.3
-1.3	1.1	-1.3	1.1	1.8
-1.7	1.1	-1	2.9	4.3
				2.4

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TABLE 2 (Page 5)

LOW	G3	T1	normal	normal	normal
TC19A_BDE_DIFF	TC20A_BDE_DIFF	TC21A_BDE_DIFF	TC22A_BDE_DIFF	TC23A_BDE_DIFF	
1.9	1.5	1	1	1	1.7
2	1.2	1.2	1	1	1.9
-1.1	-1.1	-2	1	-1	
2.5	1.1	-1.1	1	1.8	
2.3	-1.3	-1.2	1	1.1	
2.6	2.3	-1.4	1	1.5	
1.2	1	1.6	1	1.5	
2.1	2.1	-1.4	1	1.4	
-1.3	-1.2	1.2	1	1.1	
1.2	1.3	1.3	1	1.2	
1.6	1.1	1	1	1.2	
1.9	1.9	-1.1	1	1.4	
1.5	-1	1.1	1	1.2	
-1.5	-1.4	-1.4	1	-1.2	
1.7	1.3	-1	1	1.3	
-1.2	-1.1	1.4	1	1.4	
-1.2	-1.1	1.5	1	-1.2	
1.6	2.4	-1.4	1	-1.1	
2.1	1.8	-1.6	1	1	
2	2.3	-1.5	1	1.3	

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TABLE 2 (Page 6)

	Low	High	
normal	G3	Hlgh	G3
T1	T1+TIS	T1	T1
TC24A_BDE_DIFF	TC25A	TC28A_BDE_DIFF	TC29A_BDE_DIFF
-1.1	2.9	2.1	2
1.4	2	1.3	-1.1
-1	-1.1	3.4	2.9
-1.1	3.2	-1.4	2.5
2	1.3	-1	2.1
-1.8	3.8	-1.2	2.2
2.2	1.1	1.6	12
-1.6	2.4	-1.2	1.7
1.4	-1.3	1	7.1
1.6	1.5	1.4	2.5
-1.1	2.2	3.5	2
-1.3	2.3	-1.1	1.5
1.2	1.9	1.6	2.8
1.4	-1.3	2.1	1.8
-1.1	2.3	1.9	1.5
1.6	-1.2	1.1	-1
1.5	-2	3.6	2.2
-1.4	1.4	3.5	1.9
-1.4	1.6	1.5	1.3
-1.9	3.2	-1.3	1.6
			-1.2

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TABLE 2 (Page 7)

G1/2	G2	G2		
Ta	Ta/T1	Ta	INV	normal
TC31A_BDE_DIFF	TC32A_BDE_DIFF	TC33A_BDE_DIFF	TC34A_BDE_DIFF	TC35A_BDE_DIFF
1	4.4	2.5	2.3	2.3
1.1	9.1	-1.6	1.5	1.9
3.1	5.5	3.7	3.2	2.3
2.7	3.5	2	1.4	-3.6
2	2	-2.2	1.1	1.2
2.2	2.3	1.7	2.4	-1.6
4.2	2.5	-1.5	9.8	1.1
1.4	3.9	1.8	3	1.2
3.2	2.4	-1.2	6.1	1.2
6.1	3.8	1.5	2.1	1.4
2.4	2.7	2.4	1.9	1.8
1.4	3.7	1.7	2.8	1
5.5	2	2.4	2.2	3.1
2.3	3.5	2	1.9	1.7
-1.4	3.2	2	2	1.9
12	2.2	3.4	1.3	1.5
3.2	4.4	3.5	2.2	3.2
2.2	2.3	2.2	1.8	1.7
2.4	1.3	1.4	1.6	1.8
1.7	1.6	1.2	1.9	-1.7

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TABLE 2 (Page 8)

			LOW	G2/3
normal	normal	normal	Ta	T1
TC36A_BDE_DIFF	TC37A_BDE_DIFF	TC38A_BDE_DIFF	TC39A_BDE_DIFF	TC40A_BDE_DIFF
1.3	1.9	1.3	2.3	1.9
1.2	1.7	1.1	3	3.7
2.5	1.8	2.9	2.8	3.7
-1.2	-2.3	-1.6	2.4	3.2
-1.2	1.3	-1.2	6.8	1.3
-1.2	-2	-1.1	2.6	1.1
1.2	1.2	-1.1	6	4.3
1.1	-1.3	-1	3	2.1
1.5	1.2	1.2	4.7	4.5
1.6	1.7	1.4	2.2	1.3
-1.4	-1.9	1	1.3	1.6
1.1	-1.1	-1.1	3.1	2
1.9	1.8	1.6	2.5	1.6
1.8	1.7	1.6	3	2.9
1.3	1.6	1	1.7	1.6
1.3	1.9	1.3	2.6	1.9
-1.1	1	1.1	1.6	2.7
-1.3	-1.7	-1	1.4	1.6
1.3	2.3	1.2	3.3	2.8
-1.4	-2.1	-1.4	1.8	-1

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TABLE 2 (Page 9)

G2	G2	G2	G2	G2
Ta	Ta	T1	Ta	INV
TC41A BDE DIFF	TC42A BDE DIFF	TC43A BDE DIFF	TC44A BDE DIFF	TC45A BDE DIFF
1.9	2.1	2.3	2.3	2.4
2.4	2.2	4.6	3.5	1.2
3.7	3.3	3.4	3.6	3.4
1.2	2.1	1.9	3.2	-1.8
1	2.3	5	2	1.1
1.9	1.9	1.2	3.2	1.3
1.5	1.3	7	11	1.3
1.5	1.4	1.7	3.1	1.2
-1.1	1.1	4.7	6.7	1.1
2.6	2.2	2.7	2	1.7
2.3	1.6	1.9	2.8	1.2
1.5	1.6	1.8	2.5	1.2
1.7	2.1	2	2.1	2
2.6	2.1	2.4	2.8	1.5
1.4	1.6	1.6	2	2
3.1	1.4	2.6	2.8	1.3
3.1	2.3	1.6	1.4	2
2.1	1.6	1.8	2.9	1.2
2.5	1.9	1.4	2.3	1.6
1.5	1.3	1	2.6	1.3

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TABLE 2 (Page 10)

	normal	normal	normal	sectd
TC46A_BDE_DIFF	TC47A_BDE_DIFF	TC48A_BDE_DIFF		
1	1	-1.1	-1.1	-1.8
-1.2	1	-1.1		
1.7	1	3.6	1.1	
-1.4	1	-1.1	-1.2	
-1.2	1	1.1		
-1	1	-1.3	-1.6	
-1.1	1	1.1	1.3	
1.1	1	-1.2	-1.9	
-1	1	-1	1.9	
-1.2	1	1.1	-2.2	
1	1	-1.2	-2.2	
1.1	1	1.4		
-1.1	1	-1.3	-1.9	
-1.1	1	-1		
-1.1	1	1.1	3.6	
-1.1	1	-1.2	-2.3	
-1.1	1	-1.5		
-1.1	1	-1.3	-1.8	

Please renumber pages 89 to 104 of the specification as pages 106 to 121.

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REMARKS

In a March 2, 2005 telephone conference between Examiner Johannsen and Daniel N. Smith of the undersigned's office, Examiner Johannsen requested certain amendments to the specification of the above-identified application. Specifically, Examiner Johannsen requested that pages 56 to 58 of the specification be submitted as figures and that the tables on pages 83 to 88 be replaced with more legible versions.

In response, applicants attach hereto as **Exhibit A** six (6) sheets of new Figures 1-6.

Further, applicants have amended the specification to include a Brief Description of Figures section with a description of each of the figures; have deleted pages 56 to 58; and have amended pages 55 and 59 to refer to Figures 1-6. Applicants maintain that the above amendments do not create any issue of new matter and are supported, *inter alia*, on pages 55 to 59 of the application as filed.

Applicants have also amended the specification to include larger replacement tables for the tables on pages 83 to 88. Applicants maintain that these amendments also do not create any issue of new matter and are supported, *inter alia*, on pages 83 to 88 of the application as filed.

In conclusion, based on the preceding amendments to specification requested by the Examiner and the remarks, applicants respectfully request prompt issuance of the subject application for which the issue fee was paid on December 15, 2004.

If a telephone interview would be of assistance in advancing issuance of the subject application, applicants' undersigned attorney invites the Examiner to telephone him at the number

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provided below.

No fee is deemed necessary in connection with the filing of this Amendment. However, if any other fee is required, authorization is hereby given to charge the amount of such fee to Deposit Account No. 03-3125.

Respectfully submitted,


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